Clinical presentation and management of proatlas segmentation defect presenting with palatal myoclonus: case report

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Clinical presentation of craniovertebral junction disorders may range from acute catastrophic neurological deficits to insidious signs and symptoms that may mask the underlying etiology. Prompt recognition and treatment is essential to avert long-term neurological morbidity. Proatlas segmentation disorders are a rare group of developmental disorders involving the craniocervical junction. Abnormal bony segmentation leads to malformed bony structures that can in turn lead to neurological deficits through bony compression of the cervicomedullary junction. This report details a proatlas segmentation defect presenting as palatal myoclonus, a rare movement disorder. The clinical presentation, surgical management, and neuroanatomical basis for the disorder is presented. This report highlights the myriad clinical presentations of craniovertebral disorders and emphasizes a rare but treatable etiology for palatal myoclonus.

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Case Report

History and Examination

The patient was a 7-year-old boy who initially presented to the pediatric neurology service for evaluation of seizure-like episodes, as described by his parents, for the preceding 6 months. These self-limiting episodes consisted of characteristic lip smacking and tongue clicking that the child made by touching his tongue to his palate. These episodes initially lasted for 15–30 seconds and were not associated with aural, post-“ictal” phases or loss of bladder/bowel continence. He also had associated ataxia and complained of frequent headaches. In addition, the parents described behavioral problems at school, with interpersonal conflicts and emotional outbursts. His parents also described loss of hand dexterity.

After evaluation by the general pediatricians and pediatric psychiatrists, he was diagnosed with Asperger’s syndrome. On further evaluation by the pediatric neurologists, results of his initial imaging studies, including CT and MRI, were interpreted as normal. Electroencephalography and inpatient video electroencephalography monitor-
ing showed no evidence of seizure activity. He underwent genetic testing that did not reveal any underlying abnormality or syndromic condition.

A subsequent MRI study of the brain was recognized to indicate basilar invagination, and he was referred for neurosurgical evaluation. On presentation to our service, he had been experiencing these spells with increasing frequency and duration over the preceding 4–5 years. His parents described stereotypical episodes consisting of lip smacking, clucking, guttural voices, and extensor posturing with associated ataxia. Neurological examination indicated intact cranial nerve. His palate elevated symmetrically but had rhythmic clonus. Upper-extremity deep tendon reflexes showed hyperreflexia.

Neuroimaging Findings

Admission CT of the CVJ indicated proatlas segmentation abnormality with complete absence of the odontoid process (Fig. 1A). The abnormal bony segment was attached to the clivus that extended into the spinal canal, with resultant ventral cervicomedullary junction indentation (Fig. 1B, arrowhead). The MRI studies indicated a bilateral hyperintense signal within the inferior olivary nucleus (Fig. 1C, arrows). Significant ventral compression was evident (Fig. 1D). The clivus–canal angulation was significantly worse in flexion (97°) compared with extension (114°) (Fig. 1E). Tonsillar ectopia was also present. On review, these imaging findings were present in earlier imaging studies. However, radiographic progression was evident over time.

Operation

Following fiberoptic intubation and pharmacological induction of neuromuscular paralysis, intraoperative manual reduction was attempted with crown halo traction (Fig. 2A). Intraoperative CT O-arm imaging indicated irreducible ventral cervicomedullary compression and minimal improvement in cervicomedullary alignment postreduction. Hence, an anterior transpalatopharyngeal decompression of the ventral cervicomedullary junction was undertaken. Bony excision of anterior arch of the atlas, superior portion of the axis body, and midline excision of the proatlas bony segment enabled adequate ventral decompression of the medulla (Fig. 2B). Subsequently, a suboccipital C-2 pars screws, supplemented with calvarial autograft. Neurophysiological monitoring was used throughout the procedure, including the manual reduction, as previously described.

Postoperative Course

After postoperative convalescence, follow-up imaging showed adequate craniocervical decompression (Fig. 2C). Delayed dynamic radiographic imaging showed stable bony fusion (Fig. 2D). The parents reported complete resolution of the drop attacks and episodic oropharyngeal spasms. By 1-year follow-up, the child also showed significant improvement in his neurobehavioral milestones, with no further recurrences.

Discussion

Our case report highlights the neurological manifestations of CVJ disorders. Due to the anatomical juxtaposition of neural, vascular, and skeletal structures, CVJ disorders can sometimes present with unique neurological manifestations. We describe a rare presentation of symptomatic palatal tremor (SPT) as a consequence of a developmental anomaly affecting the craniocervical junction.

Palatal Tremor

Palatal tremor is a movement disorder characterized by rhythmic contractions of the soft palate. It often occurs in combination with similar rhythmic movements involving the face, larynx, and diaphragm. It has been identified by several terms, including palatal myoclonus and palatal myorhythmia, but was redefined as a palatal tremor in 1990. Palatal tremors may present either in the form of idiopathic, essential palatal tremor (EPT) or as an SPT. The essential or idiopathic form develops in the absence of any structural abnormalities or CNS lesions. The palatal tremor often occurs in combination with a self-audible clicking sound within the ear. This occurs due to pathological activation of the tensor veli palatini muscle innervated by the trigeminal cranial nerve. The resultant repetitive opening and closure of eustachian tubes is responsible for the audible clicks. Imaging findings are typically absent.

The symptomatic or secondary form occurs as a consequence of an underlying degenerative, neoplastic, vascular, structural, or demyelinating injury involving the brainstem or cerebellum. It consists of palatal contractions at a characteristic rhythmic frequency of 1–3 Hz due to involvement of the levator veli palatini muscle innervated by the glossopharyngeal cranial nerve. Ear clicks are absent and concomitant brainstem or cerebellar signs are often present due to the initial injury. Patients with SPT may also have abnormal motor learning with defective conditioning learning tasks.

Pathogenesis

The central pathogenic mechanism involves a deafferentation injury of the inferior olivary nucleus within the dentato-rubro-olivary pathway. This functional pathway, also known as the Guillain-Mollaret triangle (GMT), comprises the contralateral dentate nucleus and the ipsilateral red nucleus and inferior olivary nucleus, respectively. The dentate nucleus projects through the superior cerebellar peduncle to the contralateral red nucleus. The efferent projections comprising this dentatorubral pathway decussate in the caudal midbrain and terminate in the rostral, parvicular part of the red nucleus. Efferent projections descend in the ipsilateral central tegmental tract to the dorsal lamella of the inferior olivary nucleus. This feedback loop is completed by projections from the inferior olivary nucleus decussating fibers within the inferior cerebellar peduncle back to the dentate nucleus.

The inferior olivary nucleus is hypothesized to be a central generator with an intrinsic spontaneous activity. In palatal tremors, the deafferentation injury arises from lesions within the first 2 parts of the GMT. The site of
injury is typically ipsilateral if confined to the brainstem or contralateral if it occurs within the cerebellum. Due to the intrinsic oscillatory electrical properties of the inferior olivary nucleus, a pacemaker current is generated. This accounts for the rhythmic nature of the palatal tremor. In addition, the rhythmic pacemaker potential can also remotely modulate activity within distal spinal motor nuclei.7 Functional activation of the inferior olivary nucleus and dentate nucleus is demonstrable during the tremor periods, thereby confirming the loci.18 Tonic electromyographic activity within extremity muscles is demonstrable during palatal tremor cycles, and is most prominent in the upper rather than lower extremity and in distal versus proximal muscle groups.7 These account for the extremity and postural tremors that can be identified in patients with SPT.

The natural history of pathological and radiographic changes in relation to the onset of injury has been well described.8 Postmortem studies have helped to define the temporal profile of the associated microscopic changes.10 Neuronal hypertrophy is first demonstrable at 3 months following the lesion onset, and it peaks at 8–9 months. Following an intermediate pseudohypertrophy stage, neuronal atrophy eventually ensues. Meta-analyses have helped to delineate the temporal profile of radiographic findings in relation to the natural history of the disease.2,8 These initially include increased T2 signal intensity within 1 month of the inciting lesion, and are accompanied with a normal-sized olivary nucleus. This is followed by a phase of olivary hypertrophy due to the underlying hypertrophic olivary degeneration, with continued T2 signal hyperintensity, typically 6 months after the injury. Finally, olivary hypertrophy resolves in 3–4 years, with persistent T2 signal changes.

**Proatlas Segmentation Anomaly**

Proatlas bony segments are rare developmental anomalies of the CVJ.13 The CVJ develops by endochondral ossification. A cartilaginous framework that is initially formed is resorbed and replaced by ossification. Of the 42 somites, the craniocervical junction is formed by 4 occipital and the first 2 cervical sclerotomes.16 The first 2 occipital sclerotomes give rise to the basiocciput, and the third sclerotome develops into the jugular tubercles. The fourth occipital sclerotome or the proatlas segment normally develops into the anterior clival tubercle, the apical odontoid segment, parts of the foramen magnum margin, and the atlantal masses. The homeobox (Hox) and the paired box
The family of developmental genes regulate the embryonic development of the craniocervical junction.

Postembryonic persistence of the proatlas segment can give rise to a wide range of developmental anomalies due to the integral developmental role of the proatlas segment at the CVJ. These most commonly include a prebasilar third occipital condyle, a partial regressive occipital vertebra, or an ossiculum terminale. The proatlas segment remnant in our patient represents a developmental anomaly of the hypocentrum and centrum of the fourth occipital sclerotome. The former develops into the anterior tubercle of the clivus, and the latter forms the apical dens segment.

Proatlas segments typically present within the first 2 decades of life with clinical symptoms secondary to craniocervical compression. In the largest series to date, the most common anatomical and pathological presentation consisted of ventral craniocervical compression from abnormal bony segmentation of the clivus or the occipital condyle in up to 61% of patients. Lateral or anterolateral compression is evident in 37% and dorsal compression in 17% of the patients. Abnormal bony development of the posterior fossa leads to reduced volume, and consequently hindbrain herniation occurs in up to one-third of patients. Symptom onset typically coincides with increased physical activity and as a result of trauma in up to 55% of patients. Neurological manifestations included motor deficits in up to 72% and lower cranial nerve palsies were present in 33% of patients. Neurological symptoms may also arise from vascular compression of the vertebrobasilar system from proatlas bony remnants.

Surgical treatment is directed at decompression of the abnormal bony segment. In the event of the common ventral craniocervical compression, a transoral transpalatopharyngeal approach is necessitated. This was paired with dorsal occipitocervical fusion in almost all instances. Lateral compression is addressed via posterolateral or far lateral transcondylar approaches. Finally, dorsal compression is relieved by posterior approaches.

The diagnosis of SPT in our patient was based on the following reasons: first, the characteristic palatal tremor with the associated oropharyngeal muscular contractions in our patient is a principal feature of SPT. The facial spasms noted in our patient can be explained by coactivation of the facial nucleus with the levator veli palatini muscle. Second, coexisting brainstem and cerebellar signs, typically found in patients with SPT, were also identified in our patient in the form of ataxia, myelopathic symptoms, and pendular nystagmus. Our patient also developed extremity and postural tremor that can coexist in patients with SPT due to the underlying pacemaker activity of the inferior olivary nucleus. Third, an underlying structural cause was identified in the form of a proatlas segmentation defect with associated ventral craniocervical compression. The ventral midline compression from the prosegmentation bony anomaly probably led to dysfunction of the central tegmental tract. This also explains the bilateral nature of symptom presentation in our patient. Fourth, the clinical symptoms resolved postoperatively following surgical treatment of the underlying bony anomaly by craniovertebral decompression and stabilization. Finally, most cases...
are identifiable by T2 signal hyperintensity within the inferior olivary nucleus, as in our patient (Fig. 1C).

Conclusions

Our case report serves to emphasize the varied neurological presentations associated with CVJ disorders. Prompt recognition and diagnosis can lead to successful treatment and often neurological resolution of the underlying neurological manifestations. It also serves to highlight a rare but treatable cause for SPT in the form of a developmental CVJ disorder, with neurological recovery after surgical treatment of the underlying segmentation defect.

References


Author Contributions

Conception and design: Menezes. Acquisition of data: both authors. Analysis and interpretation of data: both authors. Drafting the article: both authors. Reviewing submitted version of manuscript: both authors. Approval of the final version of the manuscript on behalf of both authors: Menezes. Study supervision: Menezes.

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